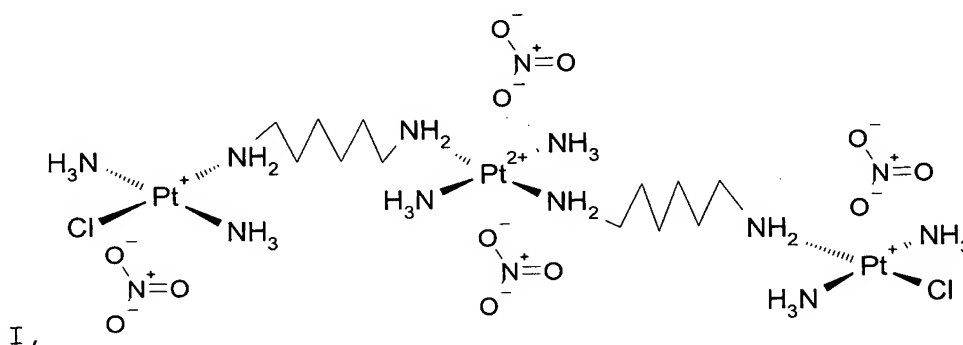


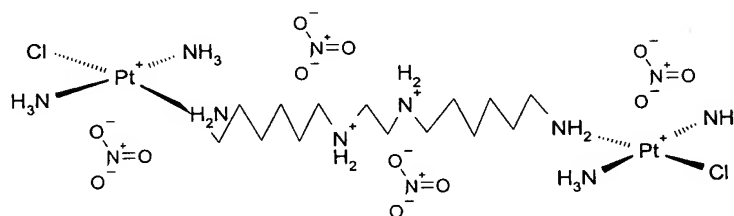
IN THE CLAIMS

1. (previously presented) Solid Lipid Nanoparticles of a platinum complex characterized by anionic ligands and ligands containing amino groups.

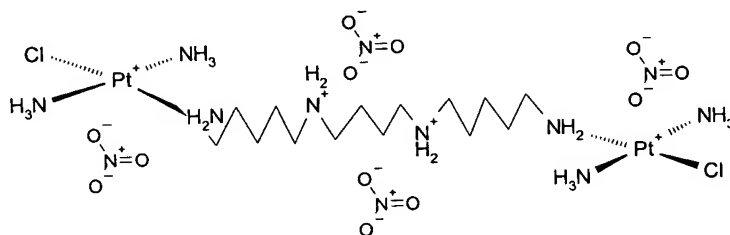
2. (currently amended) The Solid Lipid Nanoparticles of a platinum complex of according to claim 1 selected from the group consisting of trans-{bis[trans(diammine)(chloro)platinum (II) (μ -1,6-hexanediamine)]}diammineplatinum tetranitrate salt of formula I,

**Formula I**

bis{trans(diammine)(chloro)platinum(II)} μ -(1,16-diamino-7,10-diazahehexadecane-N1,N16) dinitrate salt \cdot 2HNO₃ of formula II,

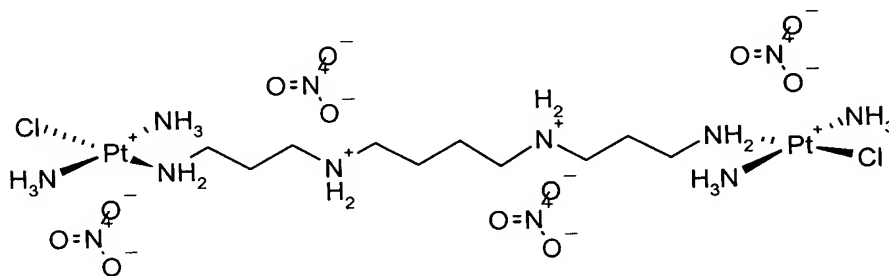
**Formula II**

bis{trans(diammine)(chloro)platinum(II)}-μ-(1,16-diamino-6,11-diazahexadecane-N1,N16) dinitrate salt- 2HNO₃ of formula III,



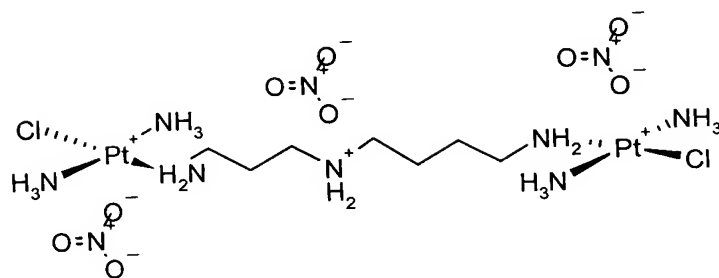
Formula III

bis{trans(diammine)(chloro)platinum(II)}-μ-(1,12-diamino-4,9-diazadodecane-N1,N12) dinitrate salt- 2HNO₃ of formula IV,



Formula IV

and bis{trans(diammine)(chloro)platinum (II)}-μ-(1,8-diamino-4-azaooctane-N¹,N⁸) dinitrate salt- HNO₃ of formula V,



Formula V.

3. (currently amended) The Solid Lipid Nanoparticles of a platinum complex of~~according to claim 1 or 2~~ obtainable by a process comprising:

- a. preparing a first microemulsion by mixing a molten lipid, a surfactant, and optionally a co-surfactant and an aqueous solution of the platinum compound~~aqueous solution~~;
- b. preparing a solution by mixing a surfactant and optionally a co-surfactant in water, heating to complete solution, preferably at the same melting temperature of the lipid used in a) and adding a co-surfactant;
- c. dispersing the microemulsion obtained in a) into the solution obtained in b) obtaining a multiple microemulsion c);
- d. dispersing the microemulsion obtained in c) in aqueous medium at a temperature ranging from 0.5°C to 4°C obtaining a dispersion of solid lipid microspheres; and
- e. washing with aqueous medium through ultrafiltration the obtained lipid microspheres obtained in d) and lyophilizing, optionally in the presence of

a bulking agent and of a cryoprotecting agent.

4. (currently amended) A process for the preparation of the Solid Lipid Nanoparticles of a platinum complex of claims 1-2, comprising:

- a. preparing a first microemulsion by mixing a molten lipid, a surfactant, and optionally a co-surfactant and an aqueous solution of the platinum complex~~compound aqueous solution~~;
- b. preparing a solution by mixing a surfactant and optionally a co-surfactant in water, heating to complete solution, preferably at the same melting temperature of the lipid used in a) and adding a co-surfactant;
- c. dispersing the microemulsion obtained in a) into the solution obtained in b) obtaining a multiple microemulsion c);
- d. dispersing the microemulsion obtained in c) in aqueous medium at a temperature ranging from 0.5°C to 4°C obtaining a dispersion of solid lipid microspheres; and
- e. washing with aqueous medium through ultrafiltration the obtained lipid microspheres obtained in d) and lyophilizing, optionally in the presence of a bulking agent and of a cryoprotecting agent.

5. (currently amended) A Pharmaceutical
pharmaceutical compositions comprising the solid-Solid lipid
Lipid nanoparticles-Nanoparticles of a platinum complex of
claims 1-3.

6. (currently amended) A method of treating a patients affected by cancer sensitive to platinum complexes,

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which comprises administering to said patients a therapeutically effective amount of the ~~solid~~ Solid lipid ~~Lipid~~ nanoparticles Nanoparticles of a platinum complex of claims 1-3.